| L | Hits | Search Text | DB | Time stamp |
|--------|--------|--|---|--------------|
| Number | 7- | | | |
| 1 | 0 | dinh-tan-thanh-\$.in. | USPAT; | 2002/07/31 |
| - | | · | US-PGPUB | 09:43 |
| 2 | 2 | dinh-tan-thanhin. | USPAT; | 2002/07/31 |
| _ | | | US-PGPUB | 09:43 |
| 3 | 0 | tremble-patrice-\$.in. | USPAT; | 2002/07/31 • |
| | | , | US-PGPUB | 09:44 |
| 4 | 1 | tremble-patricein. | USPAT; | 2002/07/31 |
| • | _ | para para para para para para para para | US-PGPUB | 09:44 |
| 5 | 11 | cunanan-crystal-\$.in. | USPAT; | 2002/07/31 |
| J | | | US-PGPUB | 09:44 |
| 6 | 0 | cunanan-crystal-m-\$.in. | USPAT; | 2002/07/31 |
| O | | Cananan Olybeal ; 4 / 2 | US-PGPUB | 09:44 |
| 7 | 11 | cunanan-crystal-min. | USPAT; | 2002/07/31 |
| ′ | | Cananan Crystar m | US-PGPUB | 09:44 |
| 8 | 0 | may-christine-\$.in. | USPAT; | 2002/07/31 |
| O | | may chilistine v.in. | US-PGPUB | 09:44 |
| 9 | 0 | may-christinein. | USPAT; | 2002/07/31 |
| 9 | , | may chiriscine .in. | US-PGPUB | 09:44 |
| 10 | 0 | ((phospholipid\$1) and (chromatagraphy or | USPAT; | 2002/07/31 |
| 10 | | tlc)).ti. | US-PGPUB | 09:45 |
| 11 | 260 | (phospholipid\$1) same (chromatagraphy or | USPAT; | 2002/07/31 |
| 11 | . 260 | tlc) | US-PGPUB | 09:46 |
| 12 | 107 | ((phospholipid\$1) same (chromatagraphy or | USPAT; | 2002/07/31 |
| 12 | 107 | tlc)) same (thin adj layer) | US-PGPUB | 09:46 |
| 13 | 6 | | USPAT; | 2002/07/31 |
| 13 | 6 | or tlc)) same (thin adj layer)) same | US-PGPUB | 09:50 |
| | | ((one adj dimensional) or (one adj way) | 05-FGF0B | 03.30 |
| | | | | |
| | 1.0 | or (one adj direction)) (((phospholipid\$1) same (chromatagraphy | USPAT; | 2002/07/31 |
| 14 | 10 | | US-PGPUB | 09:53 |
| | | or tlc)) same (thin adj layer)) and | US-FGFUB | 09.55 |
| 1.5 | 147750 | 436/71.ccls. | USPAT; | 2002/07/31 |
| 15 | 147759 | ((acetic adj acid) or ch3cooh) | US-PGPUB | 09:54 |
| 1.0 | 1.610 | (// | USPAT; | 2002/07/31 |
| 16 | 1613 | (((acetic adj acid) or ch3cooh)) same | US-PGPUB | 09:54 |
| | | ((potassium adj chloride) or kcl) | | 2002/07/31 |
| 17 | 60 | ((((acetic adj acid) or ch3cooh)) same | USPAT; | 09:54 |
| | | ((potassium adj chloride) or kcl)) same | US-PGPUB | 09:54 |
| | | (chromatography or tlc) | *************************************** | 2002/07/21 |
| 18 | 14 | (((((acetic adj acid) or ch3cooh)) same | USPAT; | 2002/07/31 |
| | | ((potassium adj chloride) or kcl)) same | US-PGPUB | 09:57 |
| | | (chromatography or tlc)) same (solvent\$1 | | |
| | | or (mobile adj phase)) | | |

WEST Search History

DATE: Wednesday, July 31, 2002

| Set Name | Hit Count | Set Name result set | | |
|--|---|------------------------|-----|--|
| DB=JPAB,EPAB,DWPI,TDBD; PLUR=YES; OP=ADJ | | | | |
| L17 | L16 and chromatography | 7 | L17 | |
| L16 | L15 and ((potassium adj chloride) or kcl) | 219 | L16 | |
| L15 | (acetic adj acid) or ch3cooh | 46629 | L15 | |
| L14 | L13 and ((one adj dimensional) or (one adj way) or (one adj direction)) | 0 | L14 | |
| L13 | phospholipid\$1 and (tlc or ((thin adj layer) adj chromatography)) | 39 | L13 | |
| L12 | wo-9950655 - \$.did. | 1 | L12 | |
| Lll | wo-9950655-\$.in. | 0 | L11 | |
| L10 | wo-0233399-\$.did. | 0 | L10 | |
| L9 | wo-02033399-\$.did. | 0 | L9 | |
| L8 | wo-2002033399-\$.did. | 0 | L8 | |
| L7 | L6 and phospholipid\$1 | 0 | L7 | |
| L6 | may-c-\$.in. | 148 | L6 | |
| L5 | L4 and phospholipid\$1 | 0 | L5 | |
| L4 | cunanan-c-\$.in. | 13 | L4 | |
| L3 | tremble-p-\$.in. | 0 | L3 | |
| L2 | L1 and phospholipid\$1 | 0 | L2 | |
| L1 | dinh-t-\$.in. | 24 | L1 | |

END OF SEARCH HISTORY

| | | E DINH TAN THANH/AU |
|-----|----------|--|
| L1 | 10 | S E1-E3 |
| L2 | 2 | S L1 AND PHOSPHOLIPID? |
| | | E TREMBLE PATRICE/AU |
| L3 | 76 | S E1-E5 |
| L4 | 1 | S L3 AND PHOSPHOLIPID? |
| | | E CUNANAN CRYSTAL M/AU |
| L5 | 16 | S E2-E3 |
| L6 | 2 | S L5 AND PHOSPHOLIPID? |
| | | E MAY CHRISTINE/AU |
| L7 | 1 | S E4 |
| F8 | 2975 | S PHOSPHOLIPID? AND (THIN LAYER CHROMATOGRAPY OR TLC) |
| L9 | 56 | S L8 AND (ONE DIMENSIONAL OR ONE WAY OR ONE DIRECTION) |
| L10 | | S L9 AND (POTASSIUM CHLORIDE OR KCL) |
| L11 | 2 | S L10 AND ACETIC ACID |
| L12 | 3 | S L10 NOT L11 |
| L13 | 5 | S L9 AND PRIMULIN |
| L14 | 3 | DUP REMOV L13 (2 DUPLICATES REMOVED) |
| L15 | 51 | S L9 NOT L10 |
| L16 | 47 | S L15 NOT L13 |
| L17 | 39 | DUP REMOV L16 (8 DUPLICATES REMOVED) |
| L18 | . 159344 | S ACETIC ACID OR CH3COOH |
| L19 | 1278 | S L18 AND (POTASSIUM CHLORIDE OR KCL) |
| L20 | | S L19 AND (CHROMATOGRAPHY OR TLC) |
| L21 | | S L20 AND (SOLVENT? OR MOBILE PHASE?) |
| L22 | 12 | DUP REMOV L21 (2 DUPLICATES REMOVED) |

=>

lysophosphatidylcholine) and three lysophospholipids (lysophosphatidylserine, lysophosphatidylethanolamine and lysophosphatidylcholine). This is achieved by simple involvement of 0.4% ammonium sulfate in silica gel H and of acetone in a developing solvent as chloroform-methanol-acetic acid-acetone-water (40:25:7:4:2). The procedure is simple and the sepn. is reproducible. The weakness of this method is the partial degrdn. of phosphatidylethanolamine to lysophosphatidylethanolamine, but a method to prevent this degrdn. is also presented.

L12 ANSWER 8 OF 12 CAPLUS COPYRIGHT 2002 ACS

Full elana Text Flatarances

AN 1992:3013 CAPLUS

DN 116:3013

TI Comparison of mobile phases for separation of **phospholipids** by **one-dimensional TLC** on preadsorbent high performance silica gel plates

AU Aloisi, Jacqueline; Fried, Bernard; Sherma, Joseph

CS Dep. Biol., Lafayette Coll., Easton, PA, 18042, USA

SO J. Liq. Chromatogr. (1991), 14(18), 3269-75 CODEN: JLCHD8; ISSN: 0148-3919

DT Journal

LA English

AB Eight solvent systems reported in the literature for the 1-dimensional TLC sepn. of phospholipids were compared under identical conditions by using high-performance preadsorbent silica gel plates. The best overall sepn. of phospholipid stds. was obtained by a single development with chloroform-methanol-water (65:25:4), and 3 other systems contg. chloroform also gave good sepns. Rf Data are tabulated for these 4 systems, and the phospholipids extd. from the digestive gland-gonad complex of Biomphalaria glabrata snails are identified.

L12 ANSWER 9 OF 12 CAPLUS COPYRIGHT 2002 ACS

Full ciang Text References

AN 1991:181515 CAPLUS

DN 114:181515

TI Comparison of mobile phases for separation and quantification of lipids by one-dimensional TLC on preadsorbent high performance silica gel plates

AU Aloisi, Jacqueline D.; Sherma, Joseph; Fried, Bernard

CS Dep. Biol., Lafayette Coll., Easton, PA, 18042, USA

SO J. Liq. Chromatogr. (1990), 13(20), 3949-61 CODEN: JLCHD8; ISSN: 0148-3919

DT Journal

LA English

Twenty-four solvent systems reported in the literature for the 1-dimensional TLC sepn. of lipids and phospholipids were compared under identical conditions by using high-performance preadsorbent silica gel plates. The best overall sepn. of mixts. of neutral lipid and phospholipid stds. and compds. extd. from the digestive gland-gonad complex of Biomphalaria glabrata snails was obtained with a system utilizing consecutive development with CHCl3-MeOH-H2O (65:25:4), CHCl3-hexane (3:1), and CCl4. The best system for quantification of neutral lipids was hexane-Et2O-HCOOH (80:20:2). Rf Data are tabulated and results discussed for all systems tested.

L12 ANSWER 10 OF 12 MEDLINE



AN 83111030 MEDLINE

DN 83111030 PubMed ID: 6822837

TI Receptor-mediated increases in phosphatidylinositol turnover in neuron-like cell lines.

L10 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2002 ACS

EUI (Oliginizi) References Text

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2002:315200 CAPLUS
ΑN
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DN 136:291337

Methods for quantitative and qualitative analyses of phospholipids using ΤI one-dimensional thin layer chromatography

Dinh, Tan Thanh; Tremble, Patrice; Cunanan, Crystal M.; Cabiling, IN Christine May

Edwards Lifesciences Corporation, USA PA

PCT Int. Appl., 23 pp. SO CODEN: PIXXD2

Patent DT

English LΑ

FAN.CNT 1

APPLICATION NO. KIND DATE PATENT NO. _____ _____ ____ WO 2001-US32023 20011012 20020425 A2 WO 2002033399 ΡI W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG PRAI US 2000-693186 20001019 Α

A highly sensitive and specific method for the detection and quantification of lipids is provided. Specifically, methods for the simultaneous detection and quantification of phospholipids extd. from mammalian tissues is described. The anal. methods provided disclose a modified one-dimensional thin-layer chromatog. technique specifically developed to rapidly and accurately detect and quantify phospholipids from mammalian cardiac tissues.

ANSWER 2 OF 2 CAPLUS COPYRIGHT 2002 ACS

Full References Text

1991:181515 CAPLUS AN

114:181515 DN

Comparison of mobile phases for separation and quantification of lipids by one-dimensional TLC on preadsorbent high performance silica gel plates

Aloisi, Jacqueline D.; Sherma, Joseph; Fried, Bernard ΑU

Dep. Biol., Lafayette Coll., Easton, PA, 18042, USA CS

J. Liq. Chromatogr. (1990), 13(20), 3949-61 SO CODEN: JLCHD8; ISSN: 0148-3919

DT Journal

English LA

Twenty-four solvent systems reported in the literature for the AB 1-dimensional TLC sepn. of lipids and phospholipids were compared under identical conditions by using high-performance preadsorbent silica gel plates. The best overall sepn. of mixts. of neutral lipid and phospholipid stds. and compds. extd. from the digestive gland-gonad complex of Biomphalaria glabrata snails was obtained with a system utilizing consecutive development with CHCl3-MeOH-H2O (65:25:4), CHCl3-hexane (3:1), and CCl4. The best system for quantification of neutral lipids was hexane-Et20-HCOOH (80:20:2). Rf Data are tabulated and results discussed for all systems tested.

7/31/02 8:06 AM

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dimensional thin-layer chromatography

- ΑU White, Thayer; Bursten, Stuart; Federighi, David; Lewis, Robert A.; Nudelman, Edward
- Cell Therapeutics, Inc., Seattle, WA, 98119, USA CS
- SO Analytical Biochemistry (1998), 258(1), 109-117 CODEN: ANBCA2; ISSN: 0003-2697
- Academic Press PB
- DT Journal
- LΑ

English An improvement of current methods is needed for simple, rapid, and precise AΒ quantification of cellular lipids, including rare species of biol. active cellular lipids, such as phosphatidic acid (PA) and diradylglycerol (DG). In addn., further anal. of hydrolyzed acyl chains from these species by methods such as gas chromatog. requires complete sepns. Methods have been developed for the quantification of neutral lipids and several phospholipids extd. from mammalian cells and sera. Lipid masses were detd. for the major classes of the neutral, nonpolar lipids, and of the phospholipids. The lipid classes were sepd. by a multistep thin-layer chromatog. (TLC) procedure in different solvent systems, a method which we have designated as multi-one-dimensional thin-layer chromatog. (MOD-TLC). Resolved lipid bands were visualized by the lipophilic dye primulin (direct yellow 59) and scanned by an automated laser-fluorescence detector. The mass of each band was detd. by comparing band intensities of unknown samples to diln. curves of authentic stds. With modifications in solvent mixts. and length of sepn. times, the majority of biol. lipids could be resolved and quantified with MOD-TLC methods. Since the detection method is nondestructive, purified lipids could then be recovered by scraping the visualized bands and extg. the lipids from the silica. The structural identities of the recovered lipids were confirmed by fast-atom bombardment and electrospray mass spectrometry. Extd. lipids were also hydrolyzed to release acylchains and acyl chain species were detd. in comparison to authentic stds. by gas chromatog. PA and DG levels in ECV.304 cells were found to be 4.6 and 3.3%, resp., of PC levels, with a PA/DG ratio of 1.4, which is in accord with published experience using other methods and different cell types. PA in human serum was detected at 0.6% of PC, indicating the sensitivity of the technique. In contrast to two-dimensional thin-layer chromatog., which allows for good resoln. of some lipid species, but cannot be used to analyze more than a single exptl. point per plate, MOD-TLC allows for direct comparative anal. of multiple samples on a single TLC plate, while still providing good resoln. for the quantification of most major classes of lipid species.

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UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
        RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,
            DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
            BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRAI US 2000-693186
                     A
                           20001019
    A highly sensitive and specific method for the detection and
    quantification of lipids is provided. Specifically, methods for the
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simultaneous detection and quantification of phospholipids extd. from mammalian tissues is described. The anal. methods provided disclose a modified one-dimensional thin-layer chromatog. technique specifically developed to rapidly and accurately detect and quantify phospholipids from mammalian cardiac tissues.

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L14 ANSWER 2 OF 3 CAPLUS COPYRIGHT 2002 ACS
   Text
    1999:641071 CAPLUS
ΑN
DN
    131:269265
    Methods of separation and detection of hydrophobic target molecules by
    multiple one-dimensional thin layer chromatography
    White, Thayer; Nudelman, Edward D.
·ΙΝ
    Cell Therapeutics, Inc., USA
PA
    PCT Int. Appl., 41.pp.
    CODEN: PIXXD2
DT
    Patent
LΑ
    English
FAN.CNT 2
    PATENT NO.
                     KIND DATE
                                         APPLICATION NO. DATE
                           -----
                     ____
                                         -----
                           19991007
                     A1
                                                          19990330
    WO 9950655
                                         WO 1999-US6803
ΡI
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W: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG AU 9934548 19991018 AU 1999-34548 A1 19990330 20011218 US 1999-465678 В1 19991217

PRAI US 6331254 US 1998-49941 A1 19980330 WO 1999-US6803 W 19990330

Methods which employ thin layer chromatog. for sepg. and detecting hydrophobic target mols. are particularly useful in sepg. biol. relevant lipids. By utilizing non-destructive detection techniques, these methods also can be adapted to further quantification or structural anal. Lipids extd. from ECV.304 cells and from pooled human serum samples were sepd. by multiple one-dimensional (MOD) TLC sequentially using chloroform-methanol-acetic acid (90:10:1, vol./vol./v), hexane-diethylether-acetone (60:40:5, vol./vol./v), hexane-diethylether (97:3, vol./vol.), and hexane (100%) as the mobile phases, all run in the same direction. The dried plates were sprayed with Primulin dye soln. and scanned by laser-excited fluorescent detection.

RE.CNT 8 THERE ARE 8 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

L14 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2002 ACS

DUPLICATE 1



1998:236214 CAPLUS ΑN

129:2268 DN

High-resolution separation and quantification of neutral lipid and TΤ phospholipid species in mammalian cells and sera by multi-one-